

Date of Report 21-01-2024
PRISCA 5.2.0.13

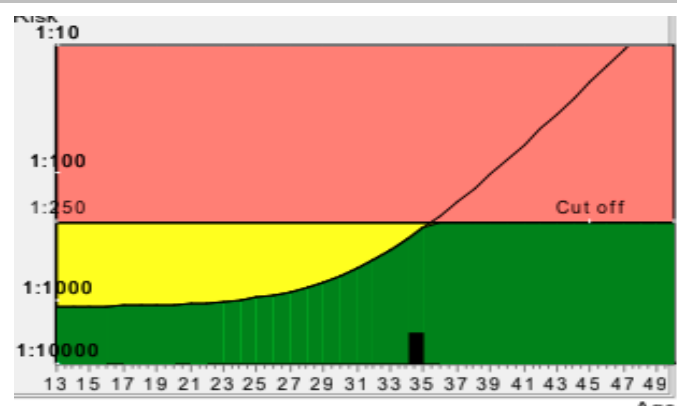
Patient Data	
Name	MRS TANVI
Birthdate	17-07-1989
Age at Sample date	34.5
Gestational age	12+4

Correction factors	
Fetuses	1 IVF
Weight in kg	54 Diabetes
Smoker	NO Origin

unknown	Previous trisomy 21	unknown
NO	Pregnancies	unknown
Asian		

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+3
PAPP-A	5.8 mIU/ml	0.93	Method	CRL (<>Robinson)
fb-hCG	63.2 ng/ml	1.65	Scan date	19-01-2024

Risks at sampling date		Ultrasound Data	
Age Risk	1:302	Crown rump length in mm	59.8
Biochemical T21 risk	1:492	Nuchal translucency MoM	0.58
Combined trisomy 21 risk	1:2678	Nasal bone	PRESENT
Trisomy 13/18 + NT	<1:10000	Sonographer	DR.INDRAJEET
		Qualifications in measuring NT	MD

Risk	Down's Syndrome Risk (Trisomy 21 Screening)
	<p>The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.</p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 2678 women with the same data, there is one woman with a trisomy 21 pregnancy and 2677 women with not affected pregnancies.</p> <p>The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that the risk calculations are statistical approaches and have no diagnostic value!</p> <p>The patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523; 1998).</p>

The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk

The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values

Risk Above Cut Off
 Risk above Age Risk
 Risk below Age risk